California Drug Recall Information

Recall Name
Hospira Recalls Magnesium Sulfate in 5% Dextrose Injection Due to Incorrect Barcode Labeling

Recall Date	Product Description	Recalling Firm	Recall Reason
3/06/15	Magnesium Sulfate in 5% Dextrose Injection, USP; 10mg/mL ***NDC # 0409-6727-23***	**Hospira, Inc.** Lake Forest, IL	*Due to confirmed customer reports of an incorrect barcode on the primary bag labeling.*
Recall Class	**Product Identification**	**Distribution**	**Affected Dates**
N/A	**Lot 42-120-JT** **Expires 1DEC2015**	**CA**, nationwide	Distributed between: October 2014 and January 2015

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm437113.htm?source=govdelivery&utm_medium=email&utm_source=govdelivery